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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(Use as many sheets as necessary)

Sheet 1 of 2

	Complete If Known
Application Number	10/539,180-Conf. #2257
Filing Date	February 28, 2005
First Named Inventor	Blas Cerda
Art Unit	1657
Examiner Name	P. C. Martin
Attorney Docket Number	NEN-23002/16

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number Number-Kind Code[2] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
	AA*	US-5,096,812	03-17-1992	Rachel et al	
	AB*	US-6,258,605	07-10-2001	Chace	
	AC*	US-6,455,321	09-24-2002	Chace	
	AD*	US-6,670,194	12-30-2003	Aebersold et al	
	AE*	US-5,629,210	05-13-1997	Hercules et al	
	AF*	US-5,719,035	02-17-1998	Rosenthal et al	

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document Country Code[3]-Number*-Kind Code[4] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages Or Relevant Figures Appear	T[5]

NON PATENT LITERATURE DOCUMENTS

Examiner Initials	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[5]
	CA	Online Mendelian Inheritance in Man (OMIM) entry for Zellweger syndrome. 2005. Accessed1 11/29/06; 13 pages	
	CB	OMIM entry for Kears-Sayre syndrome. 2006. Access online 11/29/06. 13 pages	
	CC	J.O. Sass et al., "Mutations in ACY1, the Gene Encoding Aminoacylase 1, Cause a Novel Inborn Error of Metabolism," Am. J. Hum. Genet. 2006, 78:401-409	
	CD	K.L. Peterson, D.K. Srivastava; "Functional role of a distal (3'-phosphate) group of CoA in the recombinant human liver medium-chain acyl-CoA dehydrogenase-catalysed reaction," Biochem. J 325: 751-60 , 1997.	
	CE	R. Ramsay et al., "Carnitine palmitoyltransferase in human erythrocyte membrane," Biochem. J., 1991, Vol. 275, pp. 685-88	
	CF	Definition of "metabolism," 1991. Webster's College Dictionary, Random House, page 851	
	CG	P. Rinaldo et al.; "Disorders of fatty acid transport and mitochondrial oxidation: Challenges and dilemmas of metabolic evaluation," Nov/Dec 2001;Vol. 2, No. 6, pp. 338-44	
	CH	P. Rinaldo et al., "Fatty Acid Oxidation Disorders," Annu. Rev. Physiol., 2002, 64: 477-501	
	CI	M. McCaman et al., "Fluorimetric Method for the Determination of Phenylalanine in Serum," J. Lab. Clin. Med., Vol. 59, No. 5, Aug. 1961, pp. 885, 887, 889.	
	CJ	N. Chamoles et al., "Hurler-like Phenotype: Enzymatic Diagnosis in Dried Blood Spots on Filter Paper," Clinical Chemistry, 47:12, pp. 2098-2102 , 2001.	

Examiner Signature	/Paul Martin/	Date Considered	09/11/2008

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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	CK	N. Chamoles et al., "Fabry Disease: Enzymatic Diagnosis in Dried Blood Spots on Filter Paper, Clinica Chemica Acta 308, 2001, p. 195-96	
	CL	L. Sweetman, "Newborn Screening by Tandem Mass Spectrometry (MS-MS), Clinical Chemistry, Vol. 47, No. 11, 2001, pp. 1937-38	
	CM	A. Fujimoto et al., "Quantitative Beutler Test for Newborn Mass Screening of Galactosemia Using a Fluorometric Microplate Reader, Clinical Chemistry 46:6, 2000, pp. 806-10	
Chace	CN	D. ~~Chace~~, "Rapid diagnosis of homocystinuria and other hypermethioninemias from newborns' blood spots by tandem mass spectrometry," Clinical Chemistry, 42:3, 1996, pp. 349-55	
~~—~~	~~CO~~	~~Fisher Technical Assistance website-solvent selection guide~~ no pub. date	
	CP	B. Im et al., "Bacterial Degradation of Biotin," Vol. 248, No. 22, Nov. 1973, pp. 7798-805	

Examiner Signature	/Paul Martin/	Date Considered	09/11/2008

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